

MERHAYEV, G.M.; NIKOLAYEV, A.M.

Generalized regularities of the hydrodynamics of a granular layer.
Izv.vys.ucheb.zav., khim. i khim.tekh. 8 no.2:327-330 1965.

(MIRA 28:8)

I. Yaroslavskiy tekhnologicheskiy Institut i Kazanskiy khimiko-
tekhnologicheskiy Institut Lmeni Kirova.

MIKHAYLOV, G.N., inzh.; OFMAN, A.Ya., inzh.

Improve the use of load-lifting mechanism. Bezop. truda v prom. 2
no. 6:11-12 Je '58. (MIRA 11:7)

1. Upravleniye Chelyabinskogo okruga Gosgortekhnadzora RSFSR,
(Cranes, derricks, etc.--Attachments)

OKOLOKOV, A.A., *otv. red.*; MARKIN, A.M., *otv. red.*;
BEREZOVSKIY, V.I., *red.*; DOLGUSHIN, N.I., *red.*;
KIRILLOV, I.Ye., *red.*; MIKHAYLOV, G.N., *red.*;
NEVZOROV, L.A., *red.*; NIKOLAYEVSKIY, G.M., *red.*;
ROZHDESTVENSKIY, V.A., *red.*; USHAKOV, I.N., *red.*;
KHODOV, M.P., *red.*; SHARONOV, M.S., *red.*

[Regulations for the design and safe operation of load-lifting cranes] *Pravila ustroystva i bezopasnoi ekspluatatsii gruzopod"emnykh kranov.* Moskva, Nedra, 1965. 127 p.
(MIRA 18:7)

1. Russia (1917- R.S.F.S.R.) Gosudarstvennyy komitet po nadzoru za bezopasnym vedeniyem rabot v promyshlennosti i gornomu nadzoru.

MIKHAYLOV, G.N., inzh.; OFMAN, A.Ya., inzh.

Safe operation of cranes in hot shops. Bezop.truda v prom. 2 no.9:25
S '58. (MIRA 11:9)

1.Upravleniye Chelyabinskogo okruga Gosgortekhnadzora RSFSR.
(Cranes, derricks, etc.)

USSR/Cultivated Plants. Fruits. Berries. M

Abs Jour : Ref Zhur-Biol., No 15, 1953, 63346

Author : Sapon, D. K., Ishankhanov, S. I.,
~~Mikhaylov, G. N.~~

Inst : Fruit and Berry Inst AS Uz SSR
Title : The Problem of Selecting Rootstocks for the
Saline Soils of the Golodnaya Steppe and
the Unirrigated Lands of Parkent.

Orig Pub : Tr. Flod.-yag'odn. in-ta. AN UzSSR, 1956, No 21,
143-147

Abstract : The Golodnaya Steppe and the unirrigated
lands of Parkent were investigated. The
seedlings of the following cultivated apple
strains were studied: Rosemarin, Kandil,
Sinap, Napoleon, Parkent Winter Golden, Renet

Card : 1/2

166

AUTHORS: Kabin, S. P., Mikhaylov, G. N.

48-22-3-24/30

TITLE: Dielectric Losses of Crystalline Non-Polar Polymers
(Dielektricheskiye poteri kristallicheskikh nepolyarnykh polimerov)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Seriya Fizicheskaya, 1958,
Vol. 22, Nr 3, pp. 325-328 (USSR)

ABSTRACT: In the present report the authors report on the test results of the dielectric properties on non-polar crystallizing polymers: Polyethylene and polytetrafluorethylene (ref. 1). Both polymers are typical representatives of the class of crystallizing polymers the rules of which - in a whole series of cases - differ substantially from the properties of amorphous polymers. The results may be summarized as follows:
1) Dielectric relaxation-losses can be observed both in the one as well as in the other investigated polymer. They are caused by differently constituted polar radicals which penetrate into the molecule of the polymer. 2) In the case of polyethylene, three relaxation-processes are observed at corresponding temperatures, by which it is different from amorphous polymers. 3) The amount of the losses of relaxation

Card 1/2

Dielectric Losses of Crystalline Non-Polar Polymers

48-22-3-24/30

of the mean frequency increases according to the decrease in the degree of crystallization. The relaxation-losses of the low frequency decrease. 4) In the case of polyethylene and polytetrafluorethylene, the processes of relaxation take place at the frequency of $10^4 - 10^7$ cycles per second and can be described with the same relaxation period and with one and the same activation energy, for mechanic and dielectric losses. This agreement confirms the assumption on the commonness of the molecular mechanism which are the bases for the dielectric and mechanic processes. There are 4 figures and 8 references, 7 of which are Soviet.

ASSOCIATION: Leningradskiy politekhnicheskoy institut im. M. I. Kalinina (Leningrad Polytechnical Institute imeni M. I. Kalinin). Institut vysokomolekulyarnykh soyedineniy Akademii nauk SSSR (Institute for High-Molecular Compounds AS USSR)

AVAILABLE: Library of Congress

Card 2/2
1. Polymers--Dielectric properties--Test results 2. Fluoroethylenes
--Dielectric properties--Test results 3. Polyethylene--Dielectric
properties--Test results 4. Polytetrafluorethylene--Dielectric
properties--Test results

L 23913-65 EWT(1)/EWT(m)/EEC(f)/EWA(d)/T/EWP(t)/EEC(b)-2/EWP(b) IJP(o)
JD/GG

ACCESSION NR: AP5001656

6185
S/018/64/009/012/1369/1371

AUTHOR: Artemenko, I. A.; Mykhaylov, G. O.

TITLE: Some characteristics of superconducting thin films of tin and indium

SOURCE: Ukrayins'kyy fizychnyy zhurnal, v. 9, no. 12, 1964, 1369-1371

TOPIC TAGS: superconducting indium thin film, superconducting tin film, superconductivity transition interval, indium film resistivity, tin film resistivity, superconductivity

ABSTRACT: The authors investigated superconducting thin films of tin and indium. The specimens were all of the same size (6000 Å thick, 1 x 10 mm²) and were obtained by vapor condensation of 99.999% pure metals in vacuum of 5x10⁻⁵ mm Hg on glass, mica, and quartz. Their resistances at the room and liquid helium temperatures were measured as well as the dependence of the resistances R_{243K} and R_{4.2K} on temperature and current in the film. Their ratio was by almost an order of magnitude greater in indium than in tin. The transition from

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ACCESSION NR: AP6001556

the normal to the superconducting state is in indium much sharper, the transition interval in indium being less than 0.01 degrees. This indicates that the indium films have a more homogeneous structure than the thin films. The substrate affects the characteristics of the films. Orig. art. has: 2 figures

ASSOCIATION: Institut kibernetiki AN UkrSSR, Kiev (Institute of Cybernetics AN UkrSSR)

SUBMITTED: 10Jul64

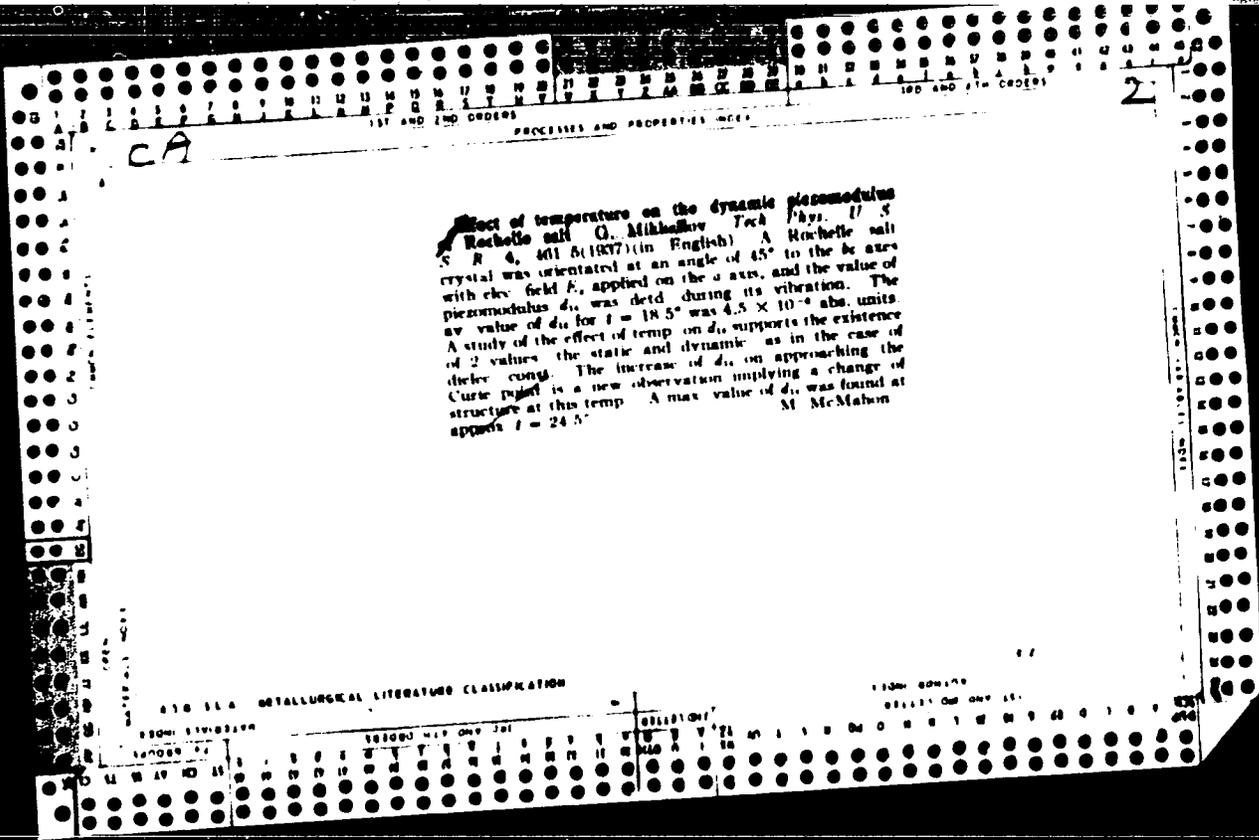
ENCL: 00

SUB CODE: SS, GP

NR REF SOV: 000

OTHER: 004

Card 2/2



Electrical methods for the measurement of the moisture
 of dispersed materials. B. P. Aleksandrov and G. Mik-
 halov. *Tech. Phys. U. S. S. R.* 5, 622-32(1938); *Chim.*
Zhur. 1939, I, 2462. —An arrangement is described for the
 detn. of capacity from measurements of the resonance
 potential which in principle is independent of the elec.
 cond. of the dielectric of the condenser. The method was
 checked experimentally by making measurements of the
 capacity of a condenser filled with water contg. varying
 concns. of electrolytes. The change in capacity and
 therefore the change in the dielec. const. were very slight
 under the given exptl. conditions. With the same app. the
 moisture content (0-8%) of a soil was detd. from measure-
 ments of the capacity of a condenser in which the soil
 served as dielectric. Even in this case the capacity was
 independent of the amt. of electrolytes contained in the soil
 at low concns. Higher concns. of electrolytes caused the in-
 crease suddenly to become worthless because of the in-
 crease in the capacity. M. G. Moore

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SA

A 53
d

4499. Influence of Electric Fields on Flow of p-Anisyanisole in Capillary Tubes. O. Mikhaylov and V. Svetkov. *J of Exp. and Theor. Phys. U.S.S.R. 2*, pp. 208-214, 1989. In Russian.—Using the capillary method it was found that a longitudinal electric field reduces and a transverse field increases the time of flow of p-anisyanisole. The effects disappear in the isotropic phase. The transverse increase occurs only if the speed of flow is not high enough to have a large orienting effect, and disappears also with increase of frequency. The authors interpret the results as being due not to a direct orienting effect of the field, but to a flow in the anisotropic liquid caused by the field. (See Abstract 1890 (1989).) D. S.

ASD 31A METALLURGICAL LITERATURE CLASSIFICATION

1st and 2nd orders PROCESSES AND PROPERTIES INDEX 3rd and 4th orders

SA

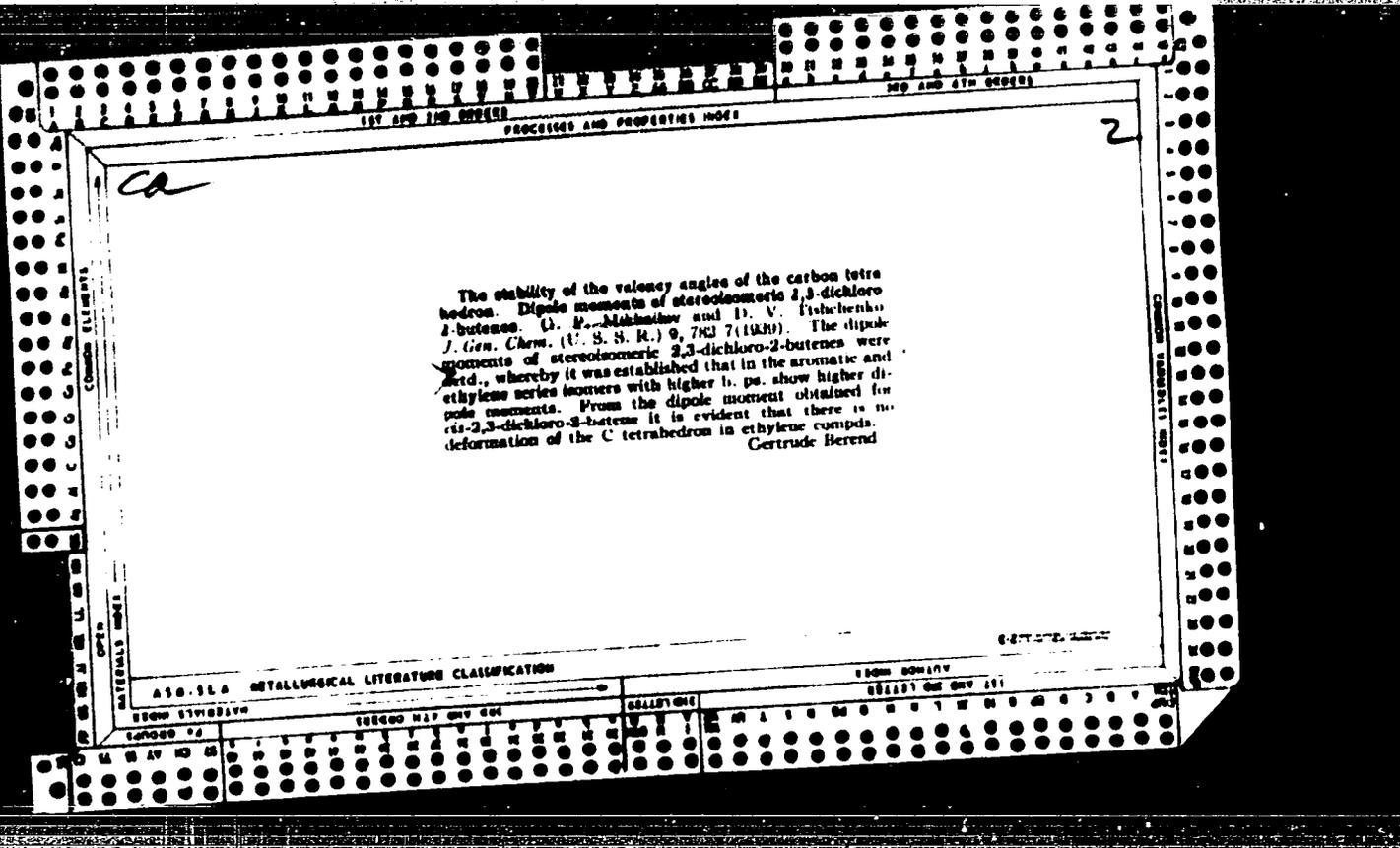
78. Action of magnetic and electric fields on flow of *p*-azoxyanisole. G. MIKHAILOV AND V. ZVETKOV. *J. of Exp. and Theor. Phys. U.S.S.R.*, 8, 8, pp. 807-801, 1939. In Russian. The paper continues previous work on the action of magnetic and electric fields on viscosity in the nematic phase (see Abstract 4490 (1939)). It is shown that the effect of a magnetic field normal to the flow direction reaches saturation for a field depending on the velocity. The viscosity of *p*-azoxyanisole when its molecules are oriented parallel to the flow but normal to the velocity gradient is found to be about 4 times the viscosity when molecules are normal to the flow but parallel to the velocity gradient. From the simultaneous action of magnetic and electric fields it is confirmed that the effect of an electric field is due to the orienting effect of the flow set up by the electric field. D. S.

A 53
V

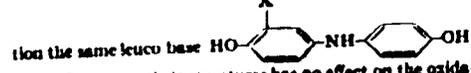
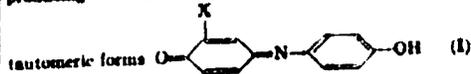
ASB 31A METALLURGICAL LITERATURE CLASSIFICATION

FROM STUBBINS 100000 100000 100000 100000

100000 100000 100000 100000



The synthesis of oxidation-reduction indicators of the indophenol series. G. Mikhalov. *Trans. Inst. Pure Chem. Reagents (U.S.S.R.)* No. 17, 45-56(1939); *Khim. Referat. Zhur.* 1940, No. 4, 30-1.—Methods for producing various indophenols are described. The



The difference in their structures has no effect on the oxidation-reduction potential if a rapid equil. is established between these forms. The individual representatives of dyes of this type are synthesized from *N*-chloroquinonimines and the subsequent condensation with the corresponding phenols. Methods for the production of the starting substance, *N*-chloroquinonimine, developed by the Inst. of Pure Chem. Reagents were described previously (C. A. 34, 3707^g). Improvements of the conditions of condensation made by the Inst. indicate that these stages of the synthesis must be carried out at a low temp. (< -2°). This increases the yield to 55% of the theoretical (based on *p*-nitrophenol) instead of 21-25%. The inner cooling used for maintaining the low temp. increases the possible charge

of *N*-chloroquinonimine used for the condensation from 5-10 mg. to 400 g. Indophenols can be purified either by dissolving and salting out or by washing with 10% NaCl. According to the modified method of Wilstätter *N*-chloroquinonimine is obtained by oxidizing *p*-aminophenol with NaOCl in a HCl medium at > 5°. Na *o*-chloroindophenolate (corresponding to II) is obtained according to Clark from the condensation of *o*-chlorophenol with quinonimine in an alk. medium. The yield is 38% of the theoretical. The tautomer I was synthesized in 72% yield in the form of the Na salt from *o*-chloroquinonimine and NaOPh in an alk. medium at > 0°. Analogous condensation of *N*, 2,6-trichloroquinonimine with NaOPh produces 75% of Na 2,6-dichloroindophenolate. Na *o*-cresolate with *N*, 2,6-trichloroquinonimine produces 98-9% of Na 2,6-dichloroindocresolate (the product contains moisture and NaCl). Na 2,6-dibromoindophenolate is obtained from the condensation of NaOPh with 2,6-dibromo-*N*-chloroquinonimine in 90% yield (based on 2,6-dibromonitrophenol). Condensation of *N*-chloroquinonimine with 1,2-C₁₀H₆(OH)SO₃Na at a temp. below -2° produces 93% of di-Na indosulfonaphthol. Owing to the presence of NaCl in the products the content of the substance varies between 60 and 80%. The compn. of the products in glass containers with ground stoppers changes very little (approx. 10% after 1.5 years). All the products possess a normal oxidation-reduction potential and are used as oxidation-reduction indicators. 2,6-Dichloroindophenol can be used as a reagent for the detn. of vitamin C (Tilman's reagent). W. R. Henn

1ST AND 2ND ORDERS PROCESSED AND PROPERTIES INDEX

BC

Constancy of the valency angles of the carbon tetrahedron.
 Dipole moments of stereoisomeric *β*-dichloro-3-butenes.
 G. P. Michailov and D. V. Tschitzchenko (*Acta Physicochim*
 / *R.S.S.*, 1960, 12, 129-136).—The dipole moments of the
 lower- and higher-boiling isomerides of *β*-dichloro-3-butene
 are 0.40 and 2.02 D, respectively, confirming that the latter
 is the *gauche* isomeride. The val. 1.61 D. is in agreement with
 that calc. on the assumption that there is no deformation of
 the valency angle. Recorded data are compared to show
 that in aromatic and ethylenic derivatives the higher-boiling
 isomeride has the greater dipole moment. F. J. G.

ASB 51A METALLURGICAL LITERATURE CLASSIFICATION

1960 51-01000
 147380

1960 51-01000
 147380

Nov 48

USSR/Engineering
Steel Alloys
Welding

"Welded Constructions From Steel With Various
Plastic Properties," G. P. Mikbaylov, A. Z.
Solomonikov, 1 P

"Avtogen Delo" No 11

From 1943-1945, authors designed and constructed
buildings from steel with increased mechanical
properties. Built welded constructions entirely
from alloyed steels and combinations of alloyed
and low-carbon steel. Carbon steel used had a
yield point of 22 kg/cm². Describes tests made on
beams using a 300-ton press preliminary to the actual
constructions. Use of alloyed steels effects a de-
crease in the weight of the construction due to in-
crease in allowable stresses.

PA 56/49742

56/49742

MIKHAYLOV, G. P.

CA

Determination of the dipole moments of copolymers. P. P. Kabanov, G. F. Mitshulov, Z. I. Novikova (Leningrad Phys. Tech. Inst.), and M. I. Kabanov. *Zhur. Tekh. Fiz.* 19, 111-18 (1949).—In extension of Debye's method of liquid mixts. of polar substances in nonpolar solvents, and of Onsager's equation (C.A. 39, 700Z) to solid copolymers of a polar and a nonpolar monomer, the dielec. const. ϵ and the dielec. losses $\tan \delta$ were detd. as functions of the temp. for copolymers of 10% methyl vinyl ketone (I) + 90% styrene and of 10% methyl methacrylate (II) + 90% styrene, the latter being the nonpolar component. Measurements with Wien's bridge at a frequency $\omega/2\pi = 10^4$ hertz gave concordant results along rising and falling temps. The behavior of the copolymers is entirely analogous to that of solns. of polar molts. in nonpolar solvents. The max. of $\tan \delta$ on the ascending branch of $\epsilon = f(T)$ corresponds to the relaxation time $\tau = 1/\omega$. The subsequent decrease is detd. by a decrease of the sp. ϵ' and by the term $\mu^2/16T$. With increasing ω , the temp. corresponding to the max. $\tan \delta$ and the inflection of ϵ moves to higher temps. T , and vice versa. Plots of $\log \tau$ detd. from the max. of $\tan \delta$, as a function of $1/T$, are linear. For the copolymer I, at 160° , $\tau = 10^{-6}$ sec., $\omega\tau = 0.06$, and at 170° , $\tau = 2 \times 10^{-6}$ sec., $\omega\tau = 0.001$; consequently, in this temp. range, $\omega\tau$ is much less than 1, and the same holds for copolymer II in the same temp. range. The mol. polarization P_s , calcd. by Onsager's equation along the descending branch $\epsilon = f(T)$, satisfies the linear relation $P_s = a + b(1/T)$, where $a =$ electronic polarization, $b = (\mu^2/h)(4\pi/9)N$. From the expl. b , the dipole moments μ per monomer unit are calcd. for I to 2.16, for II to 1.16 D. These values are close, resp., to the group dipole moment of $-\text{CO}$, 2.7-2.8, and of $-\text{CO}_2\text{Me}$, 1.0 D., from detns. in solns. Consequently, in polymers in the high-elastic state, the orientable dipoles are identical with those that undergo orientation in polar solns. ∇ These substances.

CA

Temperature dependence of the dielectric constant of copolymers. P. P. Kobeko, G. F. Mikhailov, Z. I. Novikova (Leningrad Phys. Tech. Inst.), and M. I. Kalinin. *Zhur. Tekh. Fiz.* 19, 116-19 (1949).—The relative variations, $\Delta C/C$, of the capacity of a plane condenser were detd., as a function of the temp., for polystyrene and vinyl ketone (I), and copolymers of styrene with 5, 10, 40, and 60% methyl methacrylate (II). $\Delta C/C$ decreases with rising temp. Low amts. of the polar component I or II (cf. preceding abstr.) cause this decrease to become slower, but with higher amts. the decrease becomes increasingly faster. The temp. coeff. $(1/C)\Delta C/\Delta T$ increases linearly with the content (in %) of the polar component, somewhat faster for copolymer I. Zero value of the temp. coeff. corresponds to 5% I and to 6.5% II, in agreement with the higher value of the dipole moment of I (cf. preceding abstr.). Variation of the frequency from 10^4 to 6×10^9 hertz had no effect on the temp. curve of $\Delta C/C$ for the copolymer with 10% I. The fact that, in a wide temp. range, the temp. variation of the dielec. const. due to dipole and that due to electronic polarization can be completely compensated, indicates an essential independence of the two polarizations, the dielectric polarization of the nonpolar component and the dipole polarization of the polar part. That the compensation is attained with relatively low amts. of the polar radicals is due to the greater temp. coeff. of the dipole term of the dielec. const. The changes due to thermal expansion are very small as compared with those detd. by the dipoles.

N. Thon

S A
Sect. A

Electricity

517.224.3 : 541.64
3676. Influence of orientation on dielectric losses of
polymer polymers. G. P. MAMONTOV, Zh. Tekh. Fiz.,
14, 1293-301 (No. 11, 1951) in Russian.

Strain or orientation of polymers causes maximum
values of $\tan \delta$ to occur at a higher temperature but
with unchanged magnitude. It also causes a change
in the time of relaxation of dipole-dipole losses
indicating an increase in activation energy and mole-
cular interaction. Time of relaxation corresponding
to dipole-radical losses does not change.

A. LUKASZEWICZ

MIKHAYLOV, G. P.

Effect of stretching on the dielectric losses and polarization of crystallizable polymers. G. P. Mikhailov and B. M. Feinstein (Phys. Tech. Inst. Acad. Sci. U.S.S.R., Leningrad). *Zhur. Tekh. Fiz.* 22, 759-64 (1954).—Synthetic Neoprene samples contg. 7% MgO and 5% ZnO were subjected to 200-700% stretching at -23.8 to -14.0° . The dielec. permeability under these conditions was inversely proportional to the degree of stretching. A decrease in dielec. losses was attributed to a decrease in the distribution parameter of relaxation times. The calens. showed that the values of the ΔH , ΔF , and ΔS below 150% stretching remained practically unchanged, but above it they increased considerably. The formation of cryst. phase at a higher degree of stretching was given as the explanation of these observations. A. F. Kotlov

M. A. YOUTZ
Scopus

6062

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AM

MIKHAYLOV, G.P.

USSR.

Determination of dipole moments of polymers of homogenous series of esters of methacrylic acid. G. P. Mikhaylov and L. V. Kravtsov. *Zhur. Tekh. Fiz.* 28, 1958-9 (1958). Dipole moments found for monomers and polymers of a series for 1% soln. in benzene have the same value. These data indicate (1) that the dipole moment does not change on polymerization in spite of the change in structure of the mol. as a result of the disappearance of the double bond, (2) that the effect of correlation is absent and therefore macromole. of esters of methacrylic acid in benzene soln. appear to be very flexible. And finally, dipole moment of the monomer or the monomer unit in the polymer as for a given series does not change; i.e., when a CH_2 radical is attached to a polar group also, asymmetry in a macromol. is not created. The method of Fuoss-Kirkwood with copolymerization of the method of Debye is applicable. The formula for μ with ϵ gives results 15% too low. V. N. B.

MIKHAYLOV S.P.

USSR.

The determination of the concentration of polar groups in polyethylenes by the method of dielectric losses. S. P. Mikhaylov and T. L. Borisyev. *Zhur; Tekh. Fiz.* 23, 2100-21 (1968). The presence of C=O groups in polyethylene arises from oxidation processes during the polymerization. The value of the dielectric loss, characterized by the tangent of the angle of loss, can serve as a measure of the concn. of the polar C=O groups. The concns. of the C=O groups are calcd. by use of the formula of Fuoss and Kirkwood (C.A. 35, 2046¹). The data thus obtained agree very well with data obtained by infrared spectroscopy. J. R. Leach

MIKHAYLOV, G. P.

4

155. Investigation of the dielectric losses of low-frequency relaxation in polyethylene. G. P. MIKHAYLOV, A. M. LORANOV, and B. I. SIZOV. *Journal of Applied Physics*, 1964, 35, 1553-60. The existence of low-frequency relaxation losses in polyethylene was established. These losses could be shown to be connected with the orientation of the polar groups in the crystal phase of the polyethylene. The relaxation time of the low frequency relaxation losses is slightly reduced by distension of the polyethylene specimens, whereas that of the high frequency relaxation losses is extended by specimen distension, in agreement with experience on dipole-elastic losses of amorphous polymers. SS2D24.3443

MIKHAYLOV, G. P.
USSR/Physics - Dielectrics

FD-2822

Card 1/1 Pub. 153-5/30

Author : Mikhaylov, G. P. Kabin, S. P. and Sazhin, B. I.

Title : Study of Dielectric Losses of Polyethylene

Periodical : Zhur Tekh. Fiz, 25, 590-594, 1955

Abstract : These studies are a continuation of previous works [G. P. Mikhaylov et al., ZhTF, 23, 12 (1953); ZhTF, 24, 9, (1954)] in a wider range of frequencies and temperatures. Dielectric relaxation losses were found at room temperature to be 10^{-9} sec at high frequencies, 10^{-5} sec at medium frequencies, 10^{-2} sec at low frequencies. These peculiarities of polyethylene are explained by its crystalline structure. Five USSR and 3 foreign references.

Institution :

Submitted : September 30, 1954

M. KHAYLOV, G. P.

USSR/Atomic and Molecular Physics - Physics of the Molecule, D-2

Abst Journal: Referat Zhur - Fizika, No 12, 1956, 34102

Author: Mikhaylov, G. P., Sazhin, B. I.

Institution: None

Title: Determination of Effective Dipole Moments of Molecules and of the Structural Factor of Polar Liquids and Polymers

Original Periodical: Zh. tekhn. fiziki, 1955, 25, No 10, 1696-1702

Abstract: Generalizing the polarization equation for the case of variable fields, and taking into account the deformation portion of the polarization, the authors introduce a correction to the theoretical equation for the effective dipole moment μ_{eff} . The authors calculate μ_0 , the dipole moment of the free molecules, using the Onsager equation, generalized for the case of variable fields. The values of μ , the dipole moment of the molecule in condensed matter, is calculated with the Onsager equation, which presupposes a definite molecular model. In the calculation equations for μ_{eff} , μ_0 and μ , there are included the following parameters: α , which characterizes the distribution of the relaxation time; ϵ'_m and ϵ''_m , which are the real and imaginary parts of the generalized dielectric constant. These

1 of 3

- 1 -

USSR/Academic and Molecular Physics - Physics of the Molecule, D-2

Abst Journal: Referat Zhur - Fizika, No 12, 1956, 34308

Author: Mikhaylov, G. P., Sazhin, B. I.

Institution: None

Title: Determination of Effective Dipole Moments of Molecules and of the Structural Factor of Polar Liquids and Polymers

Original Periodical: Zh. tekhn. fiziki, 1955, 25, No 10, 1696-1702

Abstract: quantities were determined from the temperature-frequency dependence of the generalized dielectric constant. In this work, the values of $\mu\sqrt{g}$, μ_0 , μ , and g (structural factor) are determined for water, for several simple alcohols, nitrobenzols, α -bromnaphthalene, normal ketones, and polymers of the homological series of ethers of metacrylic acid. The structural factor g of water and of alcohols is approximately the same and is considerably greater than unity, this indicating, in the opinion of the authors, an absence of compensation for the dipole moments during association, and a presence of close order in the parallel axes of neighboring dipoles. In the normal ketones, the structural factor is approximately unity, indicating an equal probability of directions of dipole moments of neighboring molecules.

2 of 3

- 2 -

USSR/Atomic and Molecular Physics - Physics of the Molecule, D-2

Abst Journal: Referat Zhur - Fizika, No 12, 1956, 3400

Author: Mikhaylov, G. P., Sazhin, B. I.

Institution: None

Title: Determination of Effective Dipole Moments of Molecules and of the Structural Factor of Polar Liquids and Polymers

Original Periodical: Zh. tekhn. fiziki, 1955, 25, No 10, 1696-172

Abstract: For polymers of the homological series of ethers of metacrylic acid, the structural factor is approximately the same and less than unity, this being explained by the dipole intermolecular interaction.

MIKHAYLOV, G.P.

1002

J 1061. Dielectric losses of polytetrafluoroethylene.
 G. P. MIKHAYLOV, S. P. KAMIN, and A. I. SMOLY.
 Vysok. Frekv. Fiz., 1958, 25, 2179-82. *MI*
 E.S.A. Abs., 1958, No. 853, abs. 8. In
 investigation of the dielectric characteristics of poly-
 tetrafluoroethylene dielectric relaxation losses of an
 activation energy of 19 kcal/mol were observed,
 due to relaxation processes in the amorphous parts
 of the polymer. At $t=20^{\circ}\text{C}$ the relaxation time is
 of the order of 10^{-8} sec. The relaxation time calcu-
 lated from data on the dielectric losses of the poly-
 mer agrees with the relaxation time for mechanical
 losses at ultrasonic frequencies. 882H2451.2443

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AB 233

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MIKHAILOV, G.P.; SAZHIN, B.I.; KABIN, S.P.

Dielectric losses in polyethylene. Trudy LPI no.181:201-211 '55.
(MIRA 10:1)

(Dielectrics) (Polyethylene)

Category : USSR/Atomic and Molecular Physics - Physics of high-molecular substance D-9

Abs Jour : Ref Zhur - Fizika, No 1, 1957, No 1009

Author : Kabin, S.P., Mikhaylov, G.P.

Title : On the Mechanical and Dielectric Losses of Poly-isobutylene.

Orig Pub : Zh. tekhn. fiziki, 1956, 26, No 3, 511 515

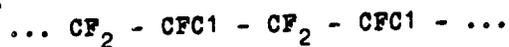
Abstract : The mechanical losses in poly-isobutylene were investigated using pulses at frequencies of 0.5 -- 3.5 Mc over a temperature range of -100 to +80°. In the interval from zero to +20° one observes a maximum of the amplitude absorption coefficient χ ; at the same time, one observes a sharp decrease in the velocity of ultrasonic waves in the polymer between -20 and +30°. The tangent of the angle of the mechanical losses has a single maximum in temperature range from -60 to +80°. Determination of the tangent of the angle of the dielectric losses of pure poly-isobutylene and of a mixture with polyethylene disclosed the presence of a maximum of dielectric losses in the same temperature interval, as the mechanical losses. The dependence of $\log f_m$ on $1/T$ (f_m is the frequency corresponding to the maximum loss) for mechanical and dielectric losses is given by one straight line, from which an activation energy of 18.6 kcal/mol was calculated. This shows that both forms of relaxation are caused by the motion of the same kinetic units of the chain.

Card : 1/1

MIKHAYLOV, P

SUBJECT USSR / PHYSICS CARD 1 / 2 PA - 1271
AUTHOR MICHAILOV, G.P., SAŽIN, B.I.
TITLE The Investigation of the Dielectric Polarization and the Losses
of Polytrifluorochlorethylene.
PERIODICAL Žurn.techn.fis, 26, fasc.8, 1723-1729 (1956)
Publ. 8 / 1956 reviewed 9 / 1956

Recently some polymers have been produced synthetically. In contrast to polytetrafluorethylene they were treated by the usual methods and have at the same time an increased resistance to heat. A typical representative of these materials is polytrifluorethylene (abbr. F-3), the molecules of which have the following structure:



As an electro-insulating material it is of considerable interest. In view of the fact that its electric properties have been but little investigated and that, besides, it is a partly crystallized polymer, the present work was carried out. On the occasion of the examination of mechanical and electric properties, a considerable influence was found to be exercised by crystal structure. Curves concerning the temperature dependence of the dielectric constant, the tg of the angle of dielectric losses, of the modulus of the displacement "G", and of the logarithmic decrement Δ of the ceasing of the mechanical oscillations of polytrifluorethylene were recorded. The dielectric losses and the polarization of F-3 are connected with molecular processes which also determine the mechanic properties of the polymer. On the occasion of the

M. KIKY, 1966, B.P.

USSR / Electricity

G

Abs Jour : Ref Zhur - Fizika N 4, 1966, No 9634

Author : Mikhaylov, S.P., Krasner, I.V.

Inst : Not given

Title : Study of the Concentration Dependence of the Relaxation Time in Copolymers by Means of the Dielectric Loss Method.

Orig Pub : Zh. tekhn. fiziki, 1966, 26, No 4, 1738-1740

Abstract : An investigation was made of the temperature dependence of the relaxation dielectric losses for copolymers of styrol with metylacrilate and styrol with metylvinylketone at various ratios of the components. All the measurements were carried out at a frequency of 1 kc. It was established that the values of $\tan \delta$ pass twice through a maximum, with the high-temperature maximum corresponding to the dipole-elastic mechanism of the losses shifting towards the region of low

Car. : 1, 2

USSR / Electricity

Abs Jour : Ref Zhur - Fizika, No 4, 1967, No 9036

Abstract : temperatures as the contents of the polar component increases, while the position of the low-temperature maximum, corresponding to the dipole-radical mechanism of losses, is independent of the composition of the copolymer. The value of $\tan \delta$ in both regions of the maximum increases with increasing content of the polar component. The value of the dielectric constant in the high-temperature loss region experiences sharp variation, and remains almost constant in the low temperature region. The results reported confirm the assumption that the dipole-elastic losses are due to the orientation of the segments of the macromolecules, while the dipole-radical losses are due to the orientation of only the lateral polar groups or monomer polar links.

Card : 2, 2

MIKHAYLOV, G.P.

USSR/ Chemistry of High-Molecular Substances

F.

Abs Jour : Referat Zhur - Khimiya, No 4, 1957, 11925

Author : Mikhaylov G.P., Borisova T.I., Dmitrochenko D.A.
Title : Dielectric Relaxation Losses in Polymethacrylate

Orig Pub : Zh. tekhn. fiziki, 1956, 26, No 9, 1924-1928

Abstract : In the frequency range of 12-80 hertz and the temperature interval of 17-120°, an investigation was made of dielectric losses of plasticized and unplasticized polymethylmethacrylate. Measurements were made by means of a Schering bridge, adapted for carrying out measurements in the above-stated frequency region. The results obtained show that frequency changes within the above-stated limits affect only the position of the region of $tg \delta$ maximum of the glassy state; a shift of the region of $tg \delta$ maximum of highly-elastic state is not observed. Addition of plasticizer affects only orientation processes of high temperature relaxation. The authors consider that losses at high temperatures are dipole-elastic, and those at low temperatures are of dipole-radical nature. Absolute values

Card 1/2

USSR/ CHEMISTRY OF HIGH-MOLECULAR SUBSTANCES

F.

Abs Jour : Referat Zhur - Khimiya, No 4, 1957, 11925

MIKHAYLOV, G. P., and BORUSOVA, T. Y.

"Dielectric measurements on polyamides," a paper presented at the 1st Congress on the Chemistry and Physics of High polymers, 26 Jan-2 Feb 57, Moscow, Fiber Research Inst.

B-3, 004, 305

MIKHAYLOV, G. P., Institute of Macromolecular Compounds, AS USSR, Leningrad

"Dielectric Losses and Polarization of Polymers," a paper submitted at the International Symposium of Macromolecular Chemistry, 9-15 Sep 1957, Prague.

MIRHAYLOV, G. P.

Mechanical and dielectric losses in polyisobutylenes.
B. P. Kabin and G. P. Mikhailov. *Soviet Phys., Tech. Phys.*
1, 493-7 (1957) (English translation). - See C.A. 50, 11081c.
B. M. R.

Handwritten marks: a checkmark, a large '2', and some illegible scribbles.

24(3)

SOV/112-58-3-3611

Translation from: Referativnyy zhurnal Elektrotehnika, 1958, Nr 3, p 9 (USSR)

AUTHOR: Mikhaylov, G. P , and Kabin, S P.

TITLE: Investigation of Dielectric Losses in Polyethylene
(Issledovaniye dielektricheskikh poter' polietilena)

PERIODICAL: Nauchno-tekhn inform byul Leningr. politekhn in-t, 1957,
Nr 4, pp 54-55

ABSTRACT: Until recently, polyethylene has been produced primarily by polymerization of ethylene under high pressure (1,000-2,000 atm) and temperatures (200°C) in the presence of oxygen acting as a catalyst. In 1955, Zigler developed a method of polymerizing ethylene under low pressures in diesel oil in the presence of $Al(C_2H_5)_3$ and $TiCl_4$. Industrial production of low-pressure polyethylene whose thermal resistance, hardness, and tensile strength exceed those of high-pressure polyethylene has been reported. It is assumed that production of low-pressure polyethylene is cheaper than that of

Card 1/2

24(3)

SOV/112-58-3-3611

Investigation of Dielectric Losses in Polyethylene

high-pressure by 30%. The present work is devoted to investigating low-pressure polyethylene produced at NIIPP & EZ (Leningrad) by a special processing method. Dielectric losses have been investigated at 50, 10^3 , 10^4 , 10^9 , and 10^{10} cps. At audio frequencies the measurements have been made within a temperature range of -100° to $+160^{\circ}$ C. It has been found that, at variance with the high-pressure polyethylene with its 3 ranges of relaxation dielectric losses, low-pressure polyethylene has only high-frequency and low-frequency relaxation losses; medium-frequency relaxation losses are absent; this is a consequence of a stronger crystallization of low-pressure polyethylene. The maximum $\text{tg}\delta \approx 3 \cdot 10^{-4}$. The $\text{tg}\delta$ of low-pressure polyethylene grows after thermal aging.

A. O. M.

Card 2/2

M.I. 11/1957
AUTHOR: BURSHTEYN, L.L. and MIKHAYLOV, G.P. PA - 2794
TITLE: Application of New Theories of Polarization to the Calculation of Effective Dipole Moments of Polar Systems. (Primeneniye novykh teoriy polarizatsii k raschetu effektivnykh dipol'nykh momentov polyarnykh sistem, Russian).
PERIODICAL: Zhurnal Tekhn. Fiz., 1957, Vol 27, Nr 4, pp 688 - 693 (U.S.S.R.)
Received: 5 / 1957 Reviewed: 6 / 1957
ABSTRACT: In the present paper the attempt is made to apply the statistical polarization theories developed by HARRIS-ALDER and PROELICH to the investigation of the dielectric polarization of a number of polar systems. The task to be performed was to obtain data on the comparison between effective dipole-moments of polar systems and solutions in which there is no interaction among polar molecules. For this purpose a method for the determination of effective dipole-moments in the case of infinite dilution was developed. A non-polar solution in infinite dilution was investigated as an example for the case of lacking interaction among polar molecules. Formulae were derived, which permit comparison of the magnitude of the effective dipole-moments and of the correlation parameters in the case of infinite dilution with the results obtained from polar liquids. The application of the new statistical theories of dielectric polarization for the computation of effective dipole-moments is demonstrated, for example, in the investigation of several representative types of the homological

Card 1/2

Application of New Theories of Polarization to the PA - 2794
Calculation of Effective Dipole Moments of Polar Systems.

ether series of metacrylic acid. According to the theory developed by HARRIS-ALDER the effective dipole-moments for these substances are computed for the case of infinite dilution. They are compared with the corresponding data for the same systems on the condition that intermolecular interaction among polar molecules is not diminated. This comparison shows that the application of the statistical theory of dielectric polarization makes it possible to form an opinion on the intermolecular interaction, which is determined by the orientation of the polar molecules with respect to each other. It may be assumed that the moments of the dipole molecules compensate one another, because the correlation parameter is considerably less than "one" and decreases with increasing μ_0 .
(5 tables and 1 citation from Slav publications)

ASSOCIATION: Institute for Polymolecular Compounds of the Academy of Science
of the U.S.S.R., Leningrad

PRESENTED BY:

SUBMITTED:

AVAILABLE: Library of Congress

Card 2/2

MIKHAYLOV G.P.

BURSHEYN, L.L. and MIKHAYLOV, G.P.

PA - 2795

Investigation of Molecular Interaction in Polymers by the Effective Dipole Moment Method. (Issledovaniye molekulyarnogo vzaimodeystviya v polimerakh metodom effektivnykh dipol'nykh momentov, Russian) Zhurnal Tekhn. Fiz., 1957, Vol 27, Nr 4, pp 694 - 701 (U.S.S.R.)

Received: 5 / 1957

Reviewed: 6 / 1957

AUTHOR:
TITLE:
PERIODICAL:
ABSTRACT:

It is shown here to what extent the investigation of dielectric polarization makes it possible to form a judgement on the molecular interaction in polymers. Even in the case of infinite dilution an interaction of the polar groups of identical chains cannot be excluded in polymers. This has an effect on the determination of the value of the effective dipole moment. Therefore it was considered appropriate to apply the statistical theories of polarization. The formula of HARRIS-ALDER was used to determine the effective dipole moment. The quantity μ (dipole moment) was computed with the help of the equation by ONZAGER. Only such polymers were investigated as contain polar groups which are more elastic than those in polyvinylchlorides or other polymers, in which the rotation of the polar groups is closely connected with the motion of the polymer chain. The effective dipole moments are determined in the case of an infinite dilution for the polymers, and the results are compared with the corresponding values for monomers in the case of infinite dilution. It appears that polymers show a smaller ratio of

ard 1/2

PA - 2795

Investigation of Molecular Interaction in Polymers by the Effective Dipole Moment Method.

μ_{eff} than monomers, and that it depends on μ_0 (dipole moment of the free molecule). This circumstance leads to suppose that an interaction exists among adjacent polar groups. In order to examine this question, such polymeric chains were investigated as follow a modification of the interaction of the polar groups. For this purpose co-polymers of methyl-methacrylate were investigated. The concentration of methyl-methacrylate varied from 24 to 80 %. The results obtained confirmed the assumption that the ratio

$\frac{\mu_{eff}}{\mu_0}$ is the determining factor of the interaction of polar groups in a macromolecular chain. Therefore investigation of dielectric polarization is actually the method to obtain data on molecular interaction in macromolecules.

(6 tables and 4 citations from Slav publications)

Card 2/2

ASSOCIATION: Institute for Highly Molecular Compounds of the Academy of Science of the U.S.S.R., Leningrad

PRESENTED BY:

SUBMITTED: 19.10.1956

AVAILABLE: Library of Congress

AUTHOR MIKHAYLOV, G.P., SAZHIN, B.I., KUPRIYANCHIK, N.N. PA 3546
TITLE: Some Peculiarities of Dielectric Losses in Polycaprolactone.
(O nekotorykh osobennostyakh dielektricheskikh poter' poli-
kaprolaktoma, Russian)
PERIODICAL: Zhurnal Tekhn. Fiz., 1957, Vol 27, Nr 5, pp 948 - 952 (U.S.S.R.)
ABSTRACT: The present work is a continuation of the two previous ones
(Zhurnal Tekhn. Fiz., 1955, Vol 25, Nr 4, p 590 and Zhurnal Tekhn.
Fiz., 1956, Vol 26, Nr 8, p 1723). The initial material was
supplied in form of little plates by the All-Union Scientific
Research Institute for Artificial Fibres. The temperature depend-
ences of the dielectric losses of hardened and tempered samples
of polycaprolactone were investigated in intervals of from 20 -
150° at frequencies of 10^3 - $5 \cdot 10^6$ Hz. In the tempered polycaprolactone
the maximum dielectric relaxation losses were found to
exist. In the case of hardened polycaprolactone a second maximum
was, in addition, noticed on the temperature curves $\text{tg } \delta$ (angle of
dielectric losses), the position of which remains the same at all
frequencies. It was possible, by means of infrared spectroscopy,
to confirm the supposition that this maximum of $\text{tg } \delta$ is connected
with an additional formation of the hydrogen compounds in the
hardened polycaprolactone within the range 30 - 40°

Card 1/2

PA - 3546

Some Peculiarities of Dielectric Losses in Polycaprolactom.

(1 illustration and 5 Slavic references)

ASSOCIATION: LPI
PRESENTED BY:
SUBMITTED: 12.11.1956
AVAILABLE: Library of Congress

Card 2/2

11-1-1000

AUTHORS Mikhaylov, G.P., Kabin, S.P. 57-9-17/40
Krylova, T.A.

TITLE On Dielectric and Mechanical Losses in Low-Pressure Polyethylene.
(O dielektricheskikh i mekhanicheskikh poteryakh polietilena nizkogo davleniya)

PERIODICAL Zhurnal Tekhn. Fiz., 1957, Vol. 27, Nr 9, pp. 2050-2055 (USSR)

ABSTRACT The results obtained by experimental investigation are given. It is shown that $\tan \delta$ of the dielectric losses within the temperature range of from -110° to $+120^{\circ}\text{C}$ and at frequencies of from 1.5 to 10 kc passes through two maximum domains. A comparison is drawn with the analogous rules for high-pressure polyethylene, and it is shown that the two types of relaxation losses in the case of low-pressure polyethylene belong to the high- and low-frequency relaxation types. Measurements of mechanical losses carried out by the ultrasonic method in dependence on temperature at a frequency of 2 kc proved the existence of only a high frequency relation. Summarizing, it is stated that the following two types of relaxation losses exist:

CARD 1/2

57-9-17/40

On Dielectric and Mechanical Losses in Low-Pressure Polyethylene.

- 1) High frequency losses reflecting the thermal motion of macromolecule components, and
- 2) losses, which reflect the thermal motion of macromolecule parts.

It is assumed that the latter are closely connected with the crystalline degree of the polymer.

There are 4 figures and 9 Slavic references.

ASSOCIATION: Leningrad Polytechnic Institute imeni M.I. Kalinin.
(Leningradskiy politekhnicheskiy institut imeni M.I. Kalinina.)

SUBMITTED: March 11, 1957.

AVAILABLE: Library of Congress.

CARD 2/2

MIKHAYLOV, G. I.

Institute of High-Molecular Compounds, USSR Acad. Sci. Leningrad

"Dielectric Losses and Relaxation of Polymers,"
paper submitted at Soviet High-Polymers, Intl. Conference, Nottingham,
UK, 21-24 July 1971.

E-3,104,641

V-60-58-4-10/18

AUTHORS: Mikhaylov, D.I., Gashin, E.I., Grebnyakova, V.I.

TITLE: Influence of the Density of Polytrifluorochloroethylene on Dielectrical Losses (Vliyanie plotnosti politriflorokhloroetilona na dielektricheskiye poteri)

PERIODICAL: Kolloidnyy zhurnal, 1959, Vol XX, Nr 4, pp 461-464

ABSTRACT: The degree of crystallization of polymers has a great effect on their physical properties. The crystallinity of a substance is defined as the volumetric or weight ratio between the crystallized and amorphous parts of this substance. It is difficult, however, to determine the exact value of crystallinity. Presently, dilatometric, optic and roentgenographic methods are used, as well as the heat capacity and the nuclear magnetic resonance. In the article, the influence of the density of polytrifluorochloroethylene (P-3) on the value of the dielectrical losses is investigated with a view to determining the degree of crystallinity. In P-3, two forms of dielectrical losses of relaxation character have been established. Both are dependent on the temperature. Figure 1 shows that at a frequency of 80 kilocycles and a temperature of 79° C a maximum of the dielectric losses is observed. The data of

Card 1 2

WV-69-58-4-10/18

Influence of the Density of Polytrifluorochloroethylene on Dielectrical
Losses

Figure 2 show that the value of the dielectric losses of P-3
is directly proportional to the quantity of the substance in
the amorphous phase. The investigation has proved that the
degree of crystallinity may be determined by measuring the di-
electric losses. The method may also be used for measuring
crystallinity in rubber, etc.
There are 2 graphs, 1 table, and 3 references, 2 of which are
Soviet and 1 English.

ASSOCIATION: Leningradskiy politekhnicheskij institut imeni V.I. Kalinina
(Leningrad Polytechnical Institute imeni V.I. Kalinin)

SUBMITTED: April 3, 1957

1. Fluoroethylenes--Density

Card 2/2

AUTHORS: Mikhaylov, G. P., Fedoseyev, G. P., 48-22-3-17/30
~~Skanavi, G. I.~~, Chmutin, M. S., Ksendzov, Ya. M.,
Matsonashvili, B. N., Kolomoitsev, F. I., Vodop'yanov, K. A.

TITLE: Discussions of Reports Submitted by: K. A. Vodop'yanov and
I. G. Vorozhtsova; K. A. Vodop'yanov and G. I. Galibina;
B. N. Matsonashvili (Preniya po dokladam: K. A. Vodop'yanova
i I. G. Vorozhtsovoy; K. A. Vodop'yanova i G. I. Galibinoy;
B. N. Matsonashvili)

PERIODICAL: Izvestiya Akademii Nauk SSSR, Seriya Fizicheskaya, 1958,
Vol. 22, Nr 3, pp. 309-310 (USSR)

ABSTRACT: G. P. Mikhaylov comments the report submitted by K. A.
Vodop'yanov and I. G. Vorozhitskaya as follows: The frequency
band is too narrow in the lecture delivered as to draw any
conclusions on the relaxation processes in mica. - G. P.
Fedoseyev says with respect to the same lecture: The lectured
conclusions are apparently scarcely convincing. The explanation
of the change-mechanism of the dielectric constant and of the
angle-tangent in mica, however, is of value. Complementary
works must be carried out, however, in order to give a more
convincing effect to the judgement on the relaxation character

Card 1/4

Discussions on Reports Submitted by: K. A. Vodop'yanov and 48-22-3-17/30
I. G. Vorozhtsova; K. A. Vodop'yanov and G. I. Galibina;
B. N. Matsonashvili

in mica. - G. I. Skanavi: Two essential contradictions exist between the works by K. A. Vodop'yanov and G. I. Galibina and the work by B. N. Matsonashvili. 1) Matsonashvili discovered relaxation maxima on the $\text{tg } \delta$ -curves of the alkaline-halogen crystals, which were not observed by Vodop'yanov and Galibina. 2) Vodop'yanov and Galibina maintain that with an increase in lattice-energy of the alkaline-halogen crystals, the $\text{tg } \delta$ decreases at room-temperature and high-frequency. Matsonashvili did not find such a correlation. The first contradiction is based on the fact that Vodop'yanov and Galibina determined the temperature dependence of $\text{tg } \delta$ on the basis of measurements at different temperatures with large temperature intervals and not in vacuum. The second contradiction may be explained by the fact that the real losses of the alkali-halogen crystals are very small at room-temperature. The losses increase rapidly, however, due to the hygroscopicity of many crystals, if no precautions were taken. M. S. Chmutin: An approximating extra-polation of the $\text{tg } \delta$ -value to high temperatures, leads - according to data by Vodop'yanov and Galibina - to a conformity with our experiments. Though

Card 2/4

Discussions of Reports Submitted by: K. A. Vodop'yanov and 48-22-7-17/30
L. G. Vorozhtsova; K. A. Vodop'yanov and G. I. Galibina,
B. N. Matsonashvili

Matsonashvili carries out his tests in vacuum, his results by extrapolation to high temperatures are higher than those of (n. M. Ksendzov: Data with smaller values of $\text{tg } \delta$, viz. the data obtained by B. N. Matsonashvili, inspire more trust.

B. N. Matsonashvili: The work-results obtained by Vodop'yanov and Galibina suffer from the fact that they were determined under atmospheric conditions. The hygroscopicity of the samples was markedly expressed in this case. Climate conditions led to the high $\text{tg } \delta$ -values I obtained. I showed in my work that the dielectric properties depend on the previous history of the sample. Therefore, only results obtained by the measurement of one and the same sample may be compared. It would be absolutely necessary to carry out a "complex" investigation of the different properties of the alkaline-halogen mono-crystals with the same samples and on the same conditions.

F. I. Kolomoitsev: It may be assumed that no fundamental contradictions exist between the experimentally obtained results which were determined in the laboratories by G. I. Skanavi and K. A. Vodop'yanov since the previous history of

Card 3/4

Discussions on Reports Submitted by: K. A. Vodop'yanov and 48-22-3-17/30
I. G. Vorozhtsova; K. A. Vodop'yanov and G. I. Galibina;
B. N. Matsonashvili

the samples may cause different results with the measuring of the $tg \delta$. - K. A. Vodop'yanov: The methods applied are the decisive factor in carrying out similar works as that by Matsonashvili and ours. The results obtained by Skarav with his method cannot deny the presence of a connection between $tg \delta$ and lattice-energy. It must be replied to G. P. Fedoseyev that it was not provided within the scope of this work to explain the practical usefulness of the thermal treatment of mica.

AVAILABLE: Library of Congress

1. Mica--Dielectric properties 2. Single crystal--dielectric properties 3. Single crystal--conductivity 4. Alkali-halogen crystals--dielectric properties

Card 4/4

AUTHORS: Mikhaylov, G. P., Lobanov, A. M. 48-22-3-22/30

TITLE: Dielectric Losses and Polarization of the Polymers (Dielektricheskiye poteri i polyarizatsiya polimerov) (Tezisy doklada) Fundamental Theses of the Report Theses (Tezisy doklada) Fundamental Theses of the Report are Given in the Article by G. P. Mikhaylov in "Progress in Chemistry" 1955, Nr 24, pp. 875 (Osnovnyye polozeniya doklada izlozhony v stat'ye G. P. Mikhaylova v "Uspekhhakh khimii" 24, 875 (1955))

PERIODICAL: Izvestiya Akademii Nauk SSSR, Seriya Fizicheskaya, 1955, Vol. 22, Nr 3, pp. 323 - 323 (USSR)

ABSTRACT: 1) The investigation of the dielectric losses and of the polarization in the wide range of both frequency and temperature showed that these properties are essentially of relaxation character. 2) According to the electric, the mechanical and other properties it was found that the polymers may be found in three physical states: in vitreous state, in highly elastic state and in viscous state. 3) The investigation of dielectric losses of the polymers in vitreous and in highly elastic state showed that these losses show a series of specific regular-

Card 1/3

48-22-3-22, 30

Dielectric Losses and Polarization of the Polymers. **Report**
Theses. Fundamental Theses of the **Report** are Given in the Article by
G. P. Mikhaylov in "Progress in Chemistry" 1955, Nr 24, pp. 875

ities. 4) A correlation is frequently observed in connection with the above explanations with the determination of the relaxation time according to the dependency of both temperature and frequency of both the dielectric and the mechanical losses. 5) Synthetic and natural polymers can be both crystalline and also amorphous, according to their structure. This is due to the corresponding properties with the dependencies of the losses and of the polarization on frequency and temperature. Such polymers as polyethylenetetrathalate, polytrifluorethylene and others are found both in crystalline and also in amorphous state. 6) Crystallization reduces considerably the dielectric losses which are correlated with the segmental thermal motion. The dielectric relaxation losses, on the other hand, which reflect the thermal motion of the monomeric terms, or of the polar radicals, change only very slightly. 7) The determination of effective dipole moments

Card 2/3

4-10-7-10

Dielectric Losses and Polarization of the Polymers. **Report**

Theses. Fundamental Theses of the **Report** are Given in the Article by G. P. Mikhaylov in "Progress in Chemistry" 1955, Nr 24, p. 1571

of the monomeric terms of the polymeric molecules, and that the theories established for monomeric molecules are applied for the calculation of the polarization of the polymers. 6) The values of the effective dipole moment were calculated for the monomeric terms of the polymers, with the prevailing chance of the value of this moment, caused by an intermolecular interaction.

ASSOCIATION: Institut vysochomolekulyarnykh soedineniy Akademiya Nauk SSSR (Institute for High Molecular Compounds, AS USSR)

AVAILABLE: Library of Congress

1. Polymers--Dielectric properties 2. Polymers--Polarization

Card 3/3

10-20-3-2, 1/1

AUTHORS: Mikhaylov, G. P., Shanavi, G. I., Tolpygo, K. B.

TITLE: Discussions on the Reports Submitted by: K. V. Filippova, I. S. Zheludev and V. M. Fridkin (Preniya po izlazu: K. V. Filippovoy, - I. S. Zhebleva i V. M. Fridkina)

PERIODICAL: Izvestiya Akademii Nauk SSSR Seriya Fizicheskaya, 1971, Vol. 22, Nr 3, p. 356 (USSR)

ABSTRACT: G. P. Mikhaylov: The results experimentally obtained by K. V. Filippova are of great interest. The tests were carried out thoroughly and accurately. The molecular explanation of the electret state, however, does not result from the report. G. I. Shanavi: The assertions by Filippova with respect to the responsibility of the dipole groups for the electret effect should be agreed with. These groups are connected with polymer formations, e. g. with segments. The orientation of the latter again causes optic anisotropy. The period of deformation of the segments is longer than that of the dipole groups at room-temperature. Contrary to the assertion by Mikhaylov, the

Card 1/2

4-22-7-29, 30

Discussions on the Reports Submitted by: K. V. Filippova, I. S. Zheludev
and V. M. Fridkin

interpretation in the actual state of work is absolutely possible and does not contradict the experimentally obtained data.

K. B. Tolpygo spoke on the lecture delivered by I. S. Zheludev and V. M. Fridkin: It seems that experimental work attained already such a stage as to allow the risk of suggesting a certain model and an energetic scheme of the electron level. The presence of basic impurity levels and levels of adhesion must apparently be assumed. The experimentally obtained results and especially the energetic distances between the $1s-1$ level and the zone of conductivity could be determined on the basis of a certain model. Concluding, attention is called to the results of discussion on the zonal theory of solid bodies which took place between the theorists of the Soviet Union.

AVAILABLE: Library of Congress

Card 2/2

1. Electrets--Properties--Theory

Investigation of Molecular Interaction in the Liquid State
Dipole Moments of Some Alcohols

The dipole moments of some alcohols in the liquid state were investigated by the method of ...
The results show that the dipole moments of these alcohols in the liquid state are significantly different from those in the gas phase. This is due to the presence of hydrogen bonds in the liquid state, which lead to a partial cancellation of the dipole moments of the individual molecules. The dipole moments of the alcohols in the liquid state are therefore smaller than those in the gas phase. The dipole moments of the alcohols in the liquid state are also affected by the temperature and the concentration of the alcohols in the liquid state. The dipole moments of the alcohols in the liquid state are therefore a function of the temperature and the concentration of the alcohols in the liquid state. The dipole moments of the alcohols in the liquid state are therefore a function of the temperature and the concentration of the alcohols in the liquid state.

Card 4/

Investigation of Molecular Interaction by the Method of Effective Dipole Moments in Some Glass-Like Polymers. 57-1-17/56

$106 \cdot 10^{18}$ erg the circle of radius corresponds to the dipole elastic
 modulus. Furthermore it is demonstrated that the character of μ_{eff}
 at a rise of temperature occurs in three stages. At first the
 effective moment nearly does not change at all only if the tem-
 perature approaches of the transition it begins to grow.
 A considerable rise of the quantity μ_{eff} is observed at T_g in the
 temperature interval of the order of magnitude of 20°C . After that
 a range of slow increase of the effective dipole moment can be
 observed. In the case of PMMA with which the temperature of gas
 formation is $+30^\circ\text{C}$ only, the third section was obtained (from
 $\mu_{eff} = \mu(t)$). The values of the effective dipole moments corres-
 ponding to the solid state of the polymers, decrease with the
 increase of the homolog number. This proves that the correlation
 increases under the influence of the groups of the side radical
 concerned. The increase of the effective dipole moment near T_g
 can be regarded as the result of a weakening of the intermolecu-
 lar interaction. Conclusively it is noted that in the glass-like
 state the motion of the polar radicals is directed preferably
 by the main chain of the polymer strand. This direction increases
 with the increase of the length of the polymer molecules which points
 out to a decrease of the effective dipole moment.

Card 3/4

Investigation of Molecular Interactions by the Method of Effective Dipole Moments on Some Crystalline-like Polymers.

There are 6 figures, 1 table and 4 references, 3 of which are Soviet.

ASSOCIATION: Institute for High-molecular Alloys A.S. USSR Leningrad (Institut vysokomolekulyarnykh soedineniy A.S. SSSR Leningrad)

SUBMITTED: June 15, 1967

AVAILABLE: Library of Congress

Card 4/4

Михайлов, Г. П.

AUTHORS: Mikhailov, G. P., Borisova, T. I.

57-1-18/30

TITLE: Investigation of Temperature-Frequency-Dependences of Dielectric Losses and Permeability of Some Polyesters of the Methacrylic Acid (Issledovaniye temperaturno-chastotnykh zavisimostey tangensa ugla dielektricheskikh poter' i pronitsayemosti nekotorykh poliefirov metakrilovoy kisloty)

PERIODICAL: Zhurnal Tekhnicheskoy Fiziki, 1958, Vol 28, Nr 1, pp. 137-142 (USSR)

ABSTRACT: This work refers to the occurrence of two types of dielectric relaxation losses demonstrated in reference 1. It is examined whether similar kinds of rules can be observed also with other representatives of the homologous series of ester of the methacrylic acid. The dependence of $\tan \delta$ and ϵ'' in the case of polymethylmethacrylate (PMMA), polyethylmethacrylate (PEMA), polybutylmethacrylate (PBMA), and polyisopropylmethacrylate (P-iso-PMA) on temperature and frequency is investigated. These investigations were carried out within the range of 12 to 10^6 C. The measuring method for $\tan \delta$ and ϵ'' , the production x) polypropylmethacrylate (PPMA)

Card 1/ 2

Investigation of Temperature-Frequency-Dependences
of Dielectric Losses and Permeability of Some
Polyesters of the Methacrylic Acid

57-1-18/30

of the polymer samples, the electrodes and other details of the experiment were the same as in reference 1. It is demonstrated that the two kinds of electrical losses occur not only with PMMA but also with other representatives of this series (PEMA, PPMA, P-iso-PMA). It was observed that the ratio between the quantities of $t_{\sigma \max}$ of dipole-elastic and dipole radical losses in the polymer series concerned changes with the increase of the number of the homolog. It was observed that the time of relaxation of the dipole radical losses in the polymer series does not change and that it shows permanently the same temperature coefficient. There are 5 figures, and 8 references, 4 of which are Slavic.

ASSOCIATION: Institute for High-molecular Alloys AN USSR Leningrad
(Institut vysokomolekulyarnykh soyedineniy AN SSSR Leningrad)
SUBMITTED: June 14, 1957
AVAILABLE: Library of Congress

Card 2/2

AUTHORS: Mikhaylov, G. P., Lobanov, A. I.

57-2-11/32

TITLE: An Investigation of the Dielectric Losses and the Permeability of Polymers **Depending on the Temperature** in the Centimeter Range of Wave-Lengths ($\lambda = 3,3$ and 10 cm) (Izucheniye dielektricheskikh poter' i pronitsayemosti polimerov v zavisimosti ot temperatury v santimetrovom diapazone dlin voln ($\lambda = 3,3$ i 10 cm)).

I. A Method for the Measurement of $\text{tg } \delta$ and ϵ' in the Temperature Interval of From -100 to 200°C at High Frequency (I. Metodika izmereniy $\text{tg } \delta$ i ϵ' v intervale temperatur ot -100 do 200°C na SVCh).

PERIODICAL: Zhurnal Tekhnicheskoy Fiziki, 1958, Vol. 28, Nr 2, pp. 267-272 (USSR).

ABSTRACT: 1.) The measurement of the temperature dependence of $\text{tg } \delta$ and ϵ' in polymers was performed at the wave-length $\lambda = 3,3$ with the aid of a gauged measuring wave guide of the type 33-И. In this occasion the method of the short-circuited wave guide was employed, where the sample was immediately fastened at the short-circuited plate. The selection of this method was due to the simple calculation formulae for this method. 2.) The dielectric measuring device ИД-1 was used for the investigations with the wave-length $\lambda = 1$ cm. This device is destined for the measure-

Card 1/2

An Investigation of the Dielectric Losses and the Permeability of Polymers in Dependence on Temperature in the Centimeter Range of Wave-Lengths ($\lambda = 3,3$ and 10 cm). 57-2-21/32

1. A Method for the Measurement of $\text{tg } \delta$ and ϵ' in the Temperature Interval of From -100 to 200°C at High Frequency.

ment according to the method of the volume-resonator of dielectrics with small losses at room temperature and $\lambda = 10$ cm. (Under 1.) were investigated: polymethylmethacrylate at 20 and 160°C , and polycaprolacton at 20 and 200°C . Under 2.) the investigations were performed in the range of from -100 to $+100^\circ\text{C}$. The experimental data and the discussion of the results follow in the next paper. The obtained data show a satisfactory accuracy in the measurements of $\text{tg } \delta$ and ϵ' depending on the temperature.

There are 3 figures, 2 tables, and 7 references, 5 of which are Slavic.

ASSOCIATION: Institute for High-Molecular Compounds AS USSR, Leningrad (Institut vyso-komplekulyarnykh soedineniy AN SSSR, Leningrad).

SUBMITTED: May 17, 1957.

AVAILABLE: Library of Congress.

Card 2/2 1. Polymers-Dielectric losses 2. Polymers-Permeability

57-2-15/32

AUTHORS: Mikhaylov, G. P., Lobov, A. A.

TITLE: An Investigation of the Dielectric Losses and the Permeability of Polymers Depending on the Frequency in the Centimeter Range of Wave-lengths ($\lambda = 1,3 \dots 10$ cm) (Issledeniye dielektricheskikh potery i pronyazhivaniya polimerov v svisi s stoyatoy temperaturoy v dlinnotolnykh volnakh ($\lambda = 1,3 \dots 10$ cm)). II. Polyethylene, polytetrafluoroethylene, Polystyrene, Polymethylmethacrylate, Polycaprolactam, Liconite (II). Polietilen, politetrafluoretilen, polistiren, polimetilmetakrilat, polikaprolaktam, likonit.

PERIODICAL: Zhurnal Tekhnicheskoy Fiziki, 1957, Vol. 27, No. 3, pp. 273-278 (USSR)

ABSTRACT: The results of the investigation of the temperature dependence of some polymers in the wave-lengths of 1,3 and 10 cm are given here. A.) Polyethylene, polytetrafluoroethylene (teflon) and polystyrene. B. δ -resonance curves ($\lambda = 1,3$ cm) are given for 1.) "high-pressure" polyethylene, 2.) "low-pressure" polyethylene, 3.) "high-pressure" polyethylene and 4.) polytetrafluoroethylene (teflon).

Card 1/4

37-15/50

An Investigation of the Dielectric Losses and the Permeability of Polymers
Depending on the Temperature in the Microwave Range of Wave-Lengths ($\lambda =$
 $= 3, 5$ and 10 cm). II. Polyethylene, Polyvinylchloride, Polystyrene,
Polymethylmethacrylate, Polymethylsiloxane (Sil-

flon) and 3,3'-polyisopropylidene "cross-linked" polystyrene
was subjected to a dielectric relaxation which caused an in-
crease in the δ value. The authors were permitted to perform
the measurements with a microwave. Curve 2 shows a maximum
which according to the authors corresponds to the losses of
the sodium-fluoride salt of the polyfrequency relaxation. (3, 2.)
and 4.) In both cases the δ value does not change with
the rise of temperature. In teflon the δ is about $1,15 \times 10^{-3}$ in
the range of $10^8 - 10^{10}$ Hz. ϵ' also remained constant
in teflon in the range of $10^8 - 10^{10}$ Hz. (3, 3.)
The value of the δ is in the range of the maximums at
room temperature and is smaller than in polyethylene.
It was difficult to determine the dielectric frequency losses from
the high-frequency losses at this level. (3, 4.) Polystyrene
an increase in the δ was observed on heating, but this can not
be called characteristic of polystyrene, as the test was per-
formed with a polystyrene of the block-polymerization. The
dielectric losses observed in polyethylene at frequencies of
the order of magnitude $10^8 - 10^{10}$ cycles prove the high flexi-

Chart 2/4

57-2-15/32

An Investigation of the Dielectric Losses and the Permeability of Polymers Depending on the Temperature in the Centimeter Range of Wave-Lengths ($\lambda = 3,3$ and 10 cm). II. Polyethylene, Polytetrafluorethylene, Polystyrene, Polymethylmethacrylate, Polycaprolactam, Ebonite

bility of the macromolecules of this polymer. B.) Polymethylmethacrylate, polycaprolactam, ebonite. 1.) The $t_g \delta$ -temperature-curve at $\lambda = 3,3$ and 10 cm for polymethylmethacrylate from -100°C shows an increase in $t_g \delta$. The displacement of the domain with a slight increase in losses towards higher temperatures in connection with a decrease in wave-length proves the relaxation-nature of the losses in polymethylmethacrylate. The ϵ'' -values at $\lambda = 3,3$ and 10 cm proved to be equal for the entire temperature range investigated. With a rise of temperature, however, ϵ'' increases, which may also be explained by the relaxation-nature of the observed losses. 2.) A similar modification of $t_g \delta$ and ϵ'' with temperature was observed in polycaprolactam and its copolymer, 95 % caprolactam and 5 % acide. The ϵ'' -values of the copolymer do not differ from those of 100 % polycaprolactam. 3.) The $t_g \delta$ -temperature- and the ϵ'' -temperature-curves for ebonite show an increase in $t_g \delta$ and ϵ'' with a rise of temperature. The

Card 3/4

57-2-15/32

An Investigation of the Dielectric Losses and the Permeability of Polymers
Depending on the Temperature in the Centimeter Range of Wave-Lengths ($\lambda =$
 $= 0,5$ and 10 cm). II. Polyethylene, Polytetrafluorethylene, Polystyrene,
Polymethylmethacrylate, Polycaprolacton, Ebonite

increase in ϵ'' with temperature also proves the relaxation-
-nature of the losses observed in ebonite. Summarizing the
authors state that with a rise of temperature $\text{tg } \delta$ due to
the occurrence of the dielectric relaxation-losses in the po-
lar polymers increases. There are 5 figures, and 10 references,
8 of which are Slavic.

ASSOCIATION: Institute of High-Molecular Compounds AS USSR, Leningrad
(Institut vysokomolekulyarnykh soyedineniy AN SSSR, Leningrad)

SUBMITTED: May 17, 1957

AVAILABLE: Library of Congress

1. Polymers-Dielectric losses 2. Polymers-Permeability

Card 4/4

IKHAYLOV, G.P.

"Untersuchung der molekularen Relaxation in Polymeren mittels der dielektrischen Methode."

Report presented at the Intl Symposium on Macromolecules, Wiesbaden, Germany, 12-27 Oct '59.

MIKHAYLOV, G.P.; SAZHIN, B.I.

Study of the dielectric losses and dielectric permittivity of
crystallizing polymers. Vysokom.soed. 1 no.1:9-16 Ja '59.
(MIRA 12:9)

1. Leningradskiy politekhnicheskoy institut im. M.I.Kalinina.
(Polymers--Electric properties)

MIKHAYLOV, G.P.; SAZHIN, B.I.

Study of the effect of polymer crystallization on dielectric losses. Vysokom.soed. 1 no.1:29-35 Ja '59. (MIRA 12:9)

1. Leningradskiy politekhnicheskoy institut im. M.I.Kalinina.
(Crystallization) (Polymers--Electric properties)

MIKHAYLOV, G.P.; BURSHTEYN, L.L.

Investigation of the dielectric polarization of methyl methacrylate
copolymers with styrene. *Viz. tver. tela* 1 no.4:632-636 '59.
(MIRA 12:6)

1. Institut vysokomolekulyarnykh soyedineniy AN SSSR, Leningrad.
(Polymers--Electric properties)
(Methacrylic acid) (Styrene)

MIKHAYLOV, G.P.; BURSHTEYN, L.L.

Study of the dipole polarization of p-chlorostyrene - isoprene
and p-chlorostyrene - styrene copolymers. *Vysokom.soed. 1*
no.12:1824-1829 D '59. (MIRA 13:5)

1. Institut vysokomolekulyarnykh soedineniy AN SSSR.
(Styrene) (Isoprene)

SOV/32-25-9-36/53

28(5)
AUTHORS:

Dmitrochenko, D. A., Lobanov, A. M., Mikhaylov, G. P.,
Shevelev, V. A.

TITLE:

Apparatus for Measuring Dielectric Losses and the Permeability
of Solid Dielectrics

PERIODICAL:

Zavodskaya laboratoriya, 1959, Vol 25, Nr 9, pp 1121-1124(USSR)

ABSTRACT:

The angles of dielectric losses $\text{tg } \delta$ and the dielectric constant ϵ within the range of decimeter waves are at present being measured according to two methods - 1) of the coaxial line, 2) of the coaxial resonator (CR) (Refs 1-5). The available constructions of (CR) do not, however, allow measurements within a wide temperature range. A (CR) has been designed, which allows measurements of the values $\text{tg } \delta$ and ϵ within a rather wide temperature range. The (CR) consists of a brass cylinder with two covers, the inner rod being soldered with silver to the inner side of the lower cover. The supports and connecting tubes of the (CR), to the generator and detector are mounted opposite each other on the side walls. The inside surfaces of the (CR) are all silver-plated and polished. The Q-factor of the (CR) attains the value 6800, with a theoretical Q-factor of 10,000. From the block diagram of the system (Fig 2) it can be seen that the (CR) is connected via an ad-

Card 1/2

SOV/32-25-9-36/53

Apparatus for Measuring Dielectric Losses and the Permeability of Solid Dielectrics

sorption-attenuator (with attenuation - 20 decibels) to a generator of type GSS-12 (frequency range 180-1000 megacycles). An amplifier 20-IM is used as indicator. Temperature was measured according to an already described method (Ref 9). The method of calculation to determine the values $\text{tg } \delta$ and ϵ is described and measurement results concerning the temperature function of these values, gathered from samples of molten quartz and polytetrafluoro ethylene (Fig 3), are quoted. The results show that the described determination method is suitable for the examination of polymer dielectrics, in which $\text{tg } \delta$ reaches magnitudes of 10^{-2} . There are 3 figures and 10 references, 5 of which are Soviet.

ASSOCIATION: Institut vysokomolekulyarnykh soyedineniy Akademii nauk BSSR (Institute for High Molecular Compounds of the Academy of Sciences, USSR)

Card 2/2

MIKHAYLOV, G.P.; BURSHTEYN, L.L.

Determining the effective dipole moments of polymers of the homologous esters of methacrylic acid in solution and in a highly elastic state. Zhur.tekh.fiz. 29 no.2:192-197 F '59.
(MIRA 12:4)

1. Institut vysokomolekulyarnykh soedineniy.
(Methacrylic acid--Dipole moments)

PHASE : FOR EMPLOYMENT

vesoyunaya konferentsii po fizike dielektrikov. 24. 1958
Fizika dielektricheskoy, tverdoy veshchiny konferentsii (Physics of Dielectrics,
Transactions of the 24 All-Union Conference on the Physics of Dielectrics)
Moscow, 1-4-56. 55 SSSR, 1960. 512 p. Errata slip inserted. 5,000 copies
printed.

Sponsoring Agency: Akademiya Nauk SSSR, Fizicheskiy Institut Iman P.M. Labeleva,
Ed. of Publishing House, Is.I. Stetsko, Vech. Ed.: I.M. Derobina, Illi-
teral Board: (Resp. Ed.) G.I. Skamri, Doctor of Physics and Mathematics
(Deceased), and E.Y. Filippov, Candidate of Physics and Mathematics.

PURPOSE: This collection of reports is intended for scientists investigating
the physics of dielectrics.

CONTENTS: The Second All-Union Conference on the Physics of Dielectrics held in
Moscow at the Fizicheskiy Institut Iman P.M. Labeleva, Physics Institute Iman
P.M. Labeleva in November 1958 was attended by representatives of the principal
scientific centers of the USSR and of several other countries. This col-
lection contains most of the reports presented at the conference and summaries
of the discussions which followed. The reports in this collection deal with
dielectric properties, losses, and polarization, and with specific technic-
capacitance of various crystals, chemical compounds, and ceramics. Photo-
electrets, ferroelectric crystals, and various radiation and irradiation ef-
fects on dielectrics are investigated. The volume contains a list of other
papers presented at the conference dealing with polarization, losses, and
breakdown of dielectrics, which were published in the journal Izvestiya AN
SSSR, seriya fizicheskaya, Nos. 4 and 5, 1959. Personalities are mentioned.
References accompany each report.

Aleksandrov, A.A., K.Ye. Likhner, and I.D. Fridlitz. Temperature Dependence
of Certain Inductive

Filktor I.D. Specific Inductive Capacitance and Dielectric Losses of Some
Ceramic Materials in Strong High-Frequency Electric Fields at High Temperatures
[Izvestiya Fiziko-khimiicheskoy Akad. Nauk SSSR, Siberian Physics and Technical
Scientific Research Institute, Tomsk].

Discussion

Dilichkov, I.I. On the Problem of the Static Specific Inductive Capacitance
of Heterogeneous Dielectrics (Vozmozhnyy mekhanizmy i teoreticheskiy
Vormozh Agricultural Institute).

Arkhangel'skiy, E.I. Dielectric Parameters of Double Liquid Systems in the
Vertical Region (Vostochny Agricultural Institute)

Isfendiyev, A.M. Anomalous Dispersion Observed in Some Dielectrics at Audio
Range (Vostochny Agricultural Institute).

Permas, Ya.M., and E.I. Labeleva. Dielectric Properties of Heterogeneous
Dielectrics at Super-High Frequencies

Discussion

Kubasov, G.B., and A.M. Labeleva. Study of ϵ' and $\tan \delta$ in Polymers as a
Function of Temperature at Super-High Frequencies (Institut Vysokichestotnykh
Vozmozh Agricultural Institute, Leningrad)

Shagin, S.M. Dielectric Characteristics ϵ' and $\tan \delta$ of Impregnated Cable
Paper in Relation to the Properties of the Components (Paper and Oil)
(Moskovskiy energeticheskiy Institut, Moscow Power Engineering Institute)

Discussion

Kashirskiy, V.Ib. Problems of the Dynamic Theory of Thermal Phenomena in
Dielectrics

Karlov, Ya.S., V.A. Kravchenko, Yu.I. Gumeny, and V.Y. Puzanov. On the
Movement of Dielectrics in an Electric Field (Leningradskiy elektrotehnicheskyy
Institut im. V.I. Il'ina, Leningrad) (Leningradskiy elektrotehnicheskyy
Institut V.I. Il'ina, Leningrad)

Dmitriyevskiy, R.A., and V.A. Shvetskiy. Use of Central Resonators for
Measuring the Dielectric Losses of Dielectrics (Leningradskiy elektrotehnicheskyy
Institut im. V.I. Il'ina, Leningrad) (Leningradskiy elektrotehnicheskyy
Institut im. V.I. Il'ina, Leningrad)

Sheludov, I.S. and V.M. Fritshteyn. Photoelectrets and the Electrolytic
Process (Izvestiya Akademiya Nauk SSSR, Moscow Institute of Geology,
Geography, Academy of Sciences USSR, Moscow).

Gubin, A.B. and V.P. Shteynman. On Charge Stability of Irganox. Kinetika
Fizicheskoy Institut Iman P.M. Labeleva, AN SSSR Moscow.

Mikhailov, G.P.

MIKHAYLOV, G.P.; EYDEL'NAFT, M.P.

Dielectric properties of a series of polyesters with aromatic nuclei
in their chains. Vysokom. soed. 2 no.2:287-294 P '60.
(MIRA 13:11)

1. Leningradskiy politekhnicheskoy institut imeni M.I.Kalinina.
(Polymers—Electric properties)

MIKHAYLOV, G.P.; EYDEL'NANT, M.P.

Temperature-frequency relationships of the angle of dielectric losses
of mixed polyesters. Vysokom. soed. 2 no.2:295-302 F '60.

(MIRA 13:11)

1. Leningradskiy politekhnicheskoy institut im.M.I.Kalinina.
(Polymers)

MIKHAYLOV, G.P.; BORISOVA, T.I.

Dielectric losses and polarization of stereoregular polymethyl
methacrylate. Vysokom. soed. 2 no.4:619-625 Ap '60.
(MIRA 13:11)

1. Institut molekulyarnykh soyedineniy AN SSSR.
(Methacrylic acid--Electric properties)

24.7800
15.8109

1144, 1147, 1145

87339

S/190/60/002/010/023/026/XX
B004/B064

AUTHORS: Mikhaylov, G. P., Eydol'nant, M. P.
TITLE: The Effect of the Polyester Structure on the Temperature Range of the Maximum of the Dielectric Losses
PERIODICAL: Vysokomolekulyarnyye soyedineniya, 1960, Vol. 2, No. 10, pp. 1548-1551

TEXT: A previous paper (Ref. 1) has dealt with the investigation of the dielectric constant and the $\tan \delta$ of the dielectric losses on various polyesters. In this connection, two types of losses were found to exist: dipole-elastic (de) and dipole-radical (dr) losses, both caused by the relaxation of the COO group. Aim of this study is to confirm the assumption that an increase of the number of methylene groups in the alcohol of the ester exerts less influence upon the temperature of the dr losses than the increase of the methylene groups in the acid. $\tan \delta$ was measured by the method described in Ref. 1 for hexamethylene terephthalate (6MT) and decamethylene terephthalate (10MT) at 1000 cps. The two substances were investigated both in elastic form, pressed from powder, and in rigid,

Card 1/4

87339

The Effect of the Polyester Structure on the
Temperature Range of the Maximum of the
Dielectric Losses

S/190/60/002/010/023/026/XX
B004/B064

crystalline form, precipitated from solution. Two maxima of $\tan \delta$ occurred as is shown in Fig. 1. The maximum at lower temperature is ascribed to the dr losses, that at higher temperature to the de losses. From the equation $\log f_{\max} = \varphi(1/T_{\max})$ (f = frequency, T = temperature) the activation energy of the dr losses was determined for 6MT to be 12, for 10MT to be 11.5 kcal/mole. A comparison with previously (Refs. 1-4) obtained data for ethylene glycol-, hexamethylene glycol-, decamethylene glycol esters of terephthalic-, adipic-, and sebacic acid led to the following conclusions: 1) the temperature range of the $\tan \delta_{\max}$ of the dielectric losses of polyesters depends linearly on the concentration of aromatic cycles in the chain of the macromolecule. There is no difference whether the concentration of the aromatic cycles changes in the alcohol or in the acid of the polyester. 2) The temperature range of the $\tan \delta_{\max}$ of the dr losses depends on the structure of the acid. For the polyesters of aromatic acids the temperature of $\tan \delta_{\max}$ is higher than for the polyesters of aliphatic acids.

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87339

The Effect of the Polyester Structure on the
Temperature Range of the Maximum of the
Dielectric Losses

S/190/60/002/010/023, 010/12
B004/B064

Legend to Fig. 1: The function $\tan \delta = \delta(t)$ for polyesters of different
crystallinity at 1000 cps. 1: 6MT pressed, 1': 6MT obtained from solution,
2: 10MT pressed, 2': 10MT obtained from solution.
There are 4 figures, 1 table, and 4 Soviet references.

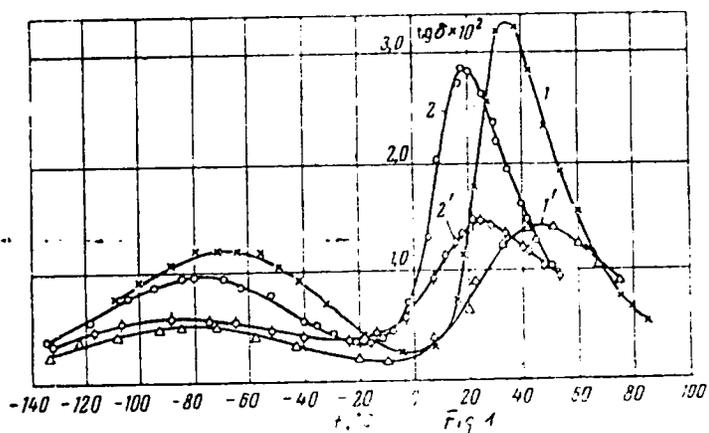
ASSOCIATION: Leningradskiy politehnicheskii institut im. M. I. Kalinina
(Leningrad Polytechnic Institute imeni M. I. Kalinin)

SUBMITTED: June 3, 1960

Card 3/4

67339

S/190/60/002/010/023/C26/XX
3004/B064



Card 4/4

87310

24.7800 1144, 1147, 1145

S/190/60/002/010/024/026, XX
B004, B064

15.8109

AUTHORS: Mikhaylov, G. P., Eydel'nant, M. P.

TITLE: Study of the Dielectric Losses of Polyethers

PERIODICAL: Vysokomolekulyarnyye soyedineniya, 1960, Vol. 2, No. 10,
pp. 1552-1556

TEXT: In continuation of previous papers (Refs. 1-3), which have dealt with the dielectric losses of polyesters, this paper discusses the investigation of the dielectric losses of polyethers: polyxylylene oxide and polyoxymethylene. Polyxylylene oxide was placed at the authors' disposal by A. V. Tunik, scientific collaborator of the IVS AN SSSR (Institute of Macromolecular Compounds of the AS USSR), polyoxymethylene by G. P. Nosov, scientific collaborator of the NIIPM (Scientific Research Institute of Plastics). $\tan \delta$ was measured by the method described in Ref. 1. The following results were obtained: Two overlapping maxima of the dielectric losses, i.e. dipole-elastic- and dipole-radical losses were observed with polyoxymethylene at -60°C . With polyxylylene oxide three maxima occur: a) at -112°C and a weaker one at -60°C (dipole-radical losses), and an

Card 1/2

87340

Study of the Dielectric Losses of Polyethers S/190/60/002/010/024/026/XX
B004/B064

intensive maximum at +10°C (dipole-elastic losses). The same laws as hold for the polyesters were observed; a) introduction of aromatic cycles increases the temperature of the $\tan \delta_{\max}$ of the dipole-elastic losses;

b) the temperature of the $\tan \delta_{\max}$ of the dipole-radical losses depends only on the mobility of the polar group (O in the case of ethers, CO in the case of esters). The dipole-radical losses are caused by the relaxation of the ether group, the dipole-elastic losses by the motion of segments of the molecular chain. There are 3 figures and 5 references: 4 Soviet and 1 German. ✓X

ASSOCIATION: Leningradskiy politekhnicheskii institut im. M. I. Kalinina
(Leningrad Polytechnic Institute imeni M. I. Kalinin)

SUBMITTED: June 3, 1960

Card 2/2

5/1/21

AUTHORS: Berezneva, G. L., Borovik, L. I., Kuznetsov, V. I.,
Mikhaylov, G. P., Nizhnik, K. I.

TITLE: Effect of Stretching on the Structure and Properties of
Films Made of Polyethylene Terephthalate. Dielectric Losses
and Electric Losses in Polyethylene Films

PERIODICAL: Vysokomolekulyarnyye soedineniya 1950, No. 1, pp. 1739-1743

TEXT: The article under consideration deals with the effect of stretching on the structure formation in thin films of polyethylene terephthalate. The authors have investigated the vitrification temperature. The authors have also investigated the conditions at which the formation of microfibrils takes place in stretched and unstretched polyethylene terephthalate films, and also the effect of stretching on the transparency and elasticity. The text of the article is written in Russian. There is a hindrance to X-ray analysis. The authors have also investigated the dielectric losses of stretched films made of polyethylene terephthalate and determined the function of the dielectric loss tangent.

Page 1/5

Effect of Stretching Upon the Structure and Properties of Films Made of Polyterephthalate. III. Study of the Losses in Face-centered Films

... in the temperature range ... of the type MJE 1 (MLYe) ... were applied to the samples ... unstretched film ... with the vitrification temperature ... at 80°C (vitrification temperature ... for 10 min at the optimum crystallization temperature ... The diagrams were constructed with ... At 1) and 2) the diagrams exhibit ... structure. In contrast therewith ... crystal structure was observed ... and no turbidity nor ... appeared. In addition ... therefrom that stretching at ... temperature ... rise to a crystallization within the primary structure, with ... being caused by spherulid structures. The ... extends the dependence of the ... of the ... structure (as found by G. P. ... and R. I. ...)

Part 1/2

Effect of Stretching Upon the Structure
and Properties of Films Made of Poly-
ethylene Terephthalate. III. Study of
Dielectric Losses in Face-centered Films

S/190/60/002/011/024/027
B004/B060

polymers submitted to mechanical pre-treatment. There are 3 figures
and 8 references: 6 Soviet, 1 US, and 1 German.

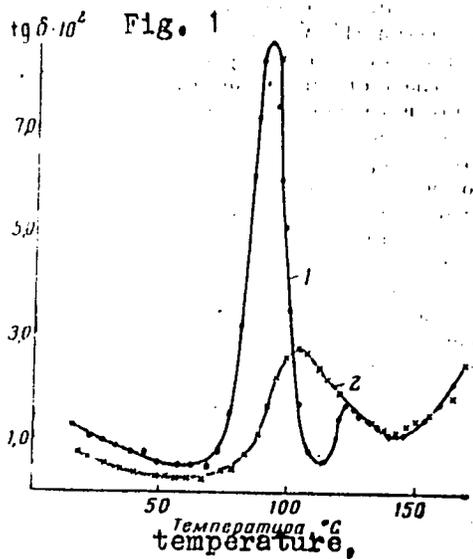
ASSOCIATION: Vsesoyuznyy nauchno-issledovatel'skiy kino-fotoinstitut
(All-Union Scientific Research Institute of Motion
Pictures and Photography). Institut vysokomolekulyarnykh
soyedineniy AN SSSR (Institute of High-molecular
Compounds of the AS USSR)

SUBMITTED: July 1, 1960

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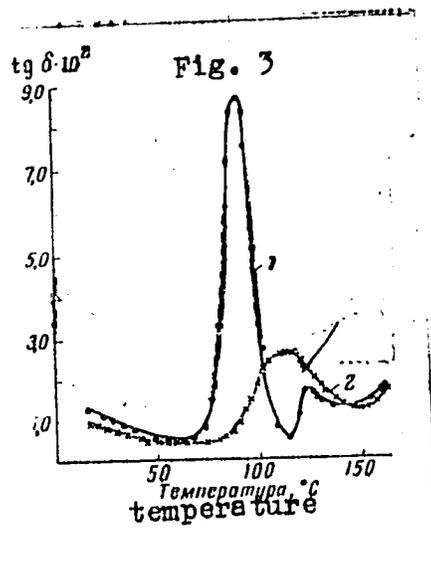


Legend to Fig.1:
 $\text{tan } \delta = \varphi(t^\circ)$ in nonoriented amorphous
polyethylene terephthalate film
($f = 1000$ cps)
1 - with temperature increase, 2 - with
temperature drop



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Legend to Fig.3:
 $\text{tan } \delta = \varphi(t^\circ)$ in face-centered polyethyl-
ene terephthalate film, stretched at 80°C ,
additionally heated to 170°C
($f = 1000 \text{ cps}$)
1 - nonoriented, amorphous sample,
2 - face-centered, heated sample.

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S/190/60/002/012/003/019

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B017/B055

AUTHORS: Mikhaylov, G. P., Borisova, T. I.

TITLE: Investigation of the Molecular Relaxation of α - and β -Chloro-substituted Polyalkyl Methacrylates by the Dielectric Loss Method

PERIODICAL: Vysokomolekulyarnye soyedineniya, 1960, Vol. 2, No. 12, pp. 1772-1778

TEXT: The authors investigated the temperature dependence of the dielectric loss ($\tan \delta$) and dielectric constant ϵ' of poly-methyl α -chloracrylate, poly-ethyl α -chloracrylate, and polypropyl α -chloracrylate, polyisopropyl α -chloracrylate and poly- β -chloro ethyl methacrylate. Measurements of $\tan \delta$ and ϵ' were carried out at frequencies of 20 - 100000 cps and temperatures of 20 - 180°C, using the apparatus described in Ref. 3. Fig. 1 shows the temperature dependence of $\tan \delta$ and ϵ' at frequencies of 20 and 400 cps for poly- α -chloro alkyl acrylate. The replacement of the CH_3 group by chlorine leads to an abrupt change in the temperature function of $\tan \delta$

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Investigation of the Molecular Relaxation of
 α - and β -Chlorosubstituted Polyalkyl
Methacrylates by the Dielectric Loss Method

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and ϵ' . The introduction of a chlorine atom causes a marked increase in dielectric loss ($\tan \delta_{\max}$). Fig. 2 represents the temperature dependence of $\tan \delta$ and ϵ' of poly- β -chloro ethyl methacrylate at 20 and 400 cps. The substitution of the CH_3 group in β -position by chlorine has no effect on the height of the dielectric loss. The frequency dependence of $\tan \delta$ and ϵ' of the polymerizates was studied in a wide temperature range and is represented graphically in Fig. 3. The occurrence of temperature-dependent dipole-elastic and dipole-radical losses was observed. The apparent activation energy of dipole relaxation in the case of polychloro alkyl methacrylates is given in a table. The determination of the dielectric relaxation of polymerizates of polyalkyl methacrylates which are substituted by chlorine in α - and β -position gave new aspects on the possibilities of preparing dielectrics with a higher dielectric constant. Measurements of ϵ' were carried out in the authors' laboratory by A. I. Artyukhov. There are 4 figures, 1 table, and 5 references: 3 Soviet, 1 US, and 1 British.

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86319

Investigation of the Molecular Relaxation of
 α - and β -Chlorosubstituted Polyalkyl
Methacrylates by the Dielectric Loss Method

S/190/60/002/012/003/019
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ASSOCIATION: Institut vysokomolekulyarnykh soyedineniy AN SSSR
(Institute of High-molecular Compounds of the Academy of
Sciences USSR)

SUBMITTED: May 12, 1960

Card 3/3

86320

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1043, 1136 only

B017/B055

AUTHORS: Mikhaylov, G. P., Borisova, T. I.

TITLE: Investigation of the Effective Dipole Moments of α - and β -Chlorine Substituted Polyalkyl Methacrylates

PERIODICAL: Vysokomolekulyarnyye soyedineniya, 1960, Vol. 2, No. 12, pp. 1779-1785

TEXT: Basing on the temperature- and frequency relationships of the generalized dielectric constant $\epsilon = \epsilon' - i\epsilon''$, the authors analyzed the relaxation-time spectrum and calculated the effective dipole moments of α - and β -chloro derivatives of polyalkyl methacrylates. The effective dipole-elastic moments $(\mu\sqrt{g})_{d.e.}$, calculated from dipole-radical loss data, are not altered by replacement of the α -CH₃ group by chlorine. The effective dipole moment as calculated from dipole-elastic relaxation, however, changes abruptly. Introduction of chlorine into the β -position produces no change. The static dipole moment $(\mu\sqrt{g})_2$ increases on intro-

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Investigation of the Effective Dipole
Moments of α - and β -Chlorine Substituted
Polyalkyl Methacrylates

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duction of chlorine into the α - or β -position of the monomer. Fig. 1 shows the relation $\epsilon'' = \varphi (\epsilon')$ for the polymers. The densities and refractive indices of several polyalkyl methacrylates are listed in Table 1. Fig. 2 represents the temperature function of the parameters of relaxation. The effective dipole moments of several polyalkyl methacrylates are given in Table 2. The dipole moments of glassy polymers increase slightly with temperature. The data obtained were compared with the corresponding data for the homologous series of polyesters of methacrylic acid. The influence of a second polar group on the relaxation properties of the polymers as a function of the position of this dipole was discussed. There are 2 figures, 2 tables, and 6 references: 4 Soviet.

ASSOCIATION: Institut vysokomolekulyarnykh soyedineniy AN SSSR
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Sciences USSR)

SUBMITTED: May 12, 1960

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